AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A dosage form of dalbavancin for parenteral use comprising:

a sterile, stable, particle-free dalbavancin powder suitable for reconstitution with a pharmaceutically acceptable vehicle; and

a stabilizer,

wherein the dosage form is at a pH of about 3-5.

2. (Original) The dosage form of claim 1, wherein the stabilizer comprises sugar.

3. (Original) The dosage form of claim 2, wherein the sugar is selected from the group consisting of mannitol, lactose, sucrose, sorbitol, glycerol, cellulose, trehalose, maltose, dextrose, and combinations thereof.

4. (Original) The dosage form of claim 1, wherein the stabilizer is mannitol.

5. (Original) The dosage form of claim 4, wherein the weight ratio of mannitol: dalbavancin is 1:2.

6. (Original) The dosage form of claim 1, wherein the stabilizer is lactose.

7. (Original) The dosage form of claim 1, wherein the stabilizer is a mixture of mannitol and lactose.

8. (Original) The dosage form of claim 7, wherein the weight ratio of mannitol:lactose:dalbavancin is 1:1:4.

9. (Original) The dosage form of claim 8, wherein the pH is 4.5.

10. (Original) The dosage form of claim 1, wherein the pH is 3.5.

11. (Original) The dosage form of claim 1, wherein the pH is 4.5.

12-66. (Canceled)